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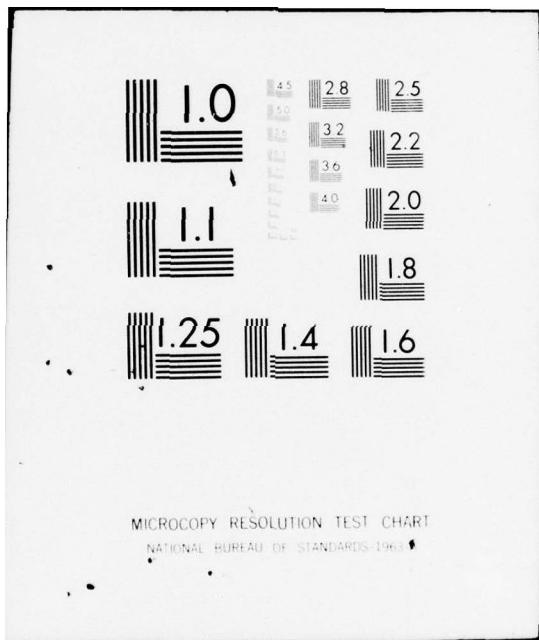
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August 1977

**STUDY OF EROSION MECHANISMS
OF ENGINEERING CERAMICS**

Sixth Interim Technical Report
April 1 to July 31, 1977

Prepared Under Contract N00014-73-C-0401
NR0-32-542

by
M. E. Gulden

for

**OFFICE OF NAVAL RESEARCH
DEPARTMENT OF THE NAVY**

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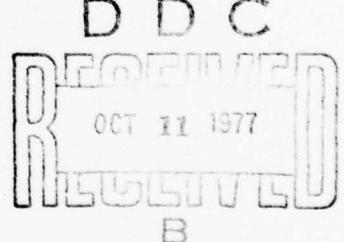
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FOREWORD

This report covers the experimental work performed by Solar from April 1, 1977 to July 31, 1977 on Contract N00014-73-C-0401 sponsored by the Office of Naval Research, Department of the Navy. Relevant observations from previous work on this contract are also included. Dr. A. Diness of the Metallurgy Program is monitoring this contract. The work was performed by Mary Ellen Gulden, Senior Research Engineer, as Principal Investigator under the direction of Dr. Arthur G. Metcalfe, Associate Director of Research. The testing was performed by R. Bradley Domes, Senior Research Technician.

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ABSTRACT

Four "engineering" ceramics were subjected to impact (single particle) and erosion (multiple impacts) under conditions which simulate a natural dust environment in the subsonic velocity regime. The target materials are hot pressed Si_3N_4 , reaction bonded Si_3N_4 , glass bonded Al_2O_3 and hot pressed MgF_2 . Tests were performed with 6 narrow size ranges of natural quartz between 10 and 385μ average, and 5 velocities for each particle size. Hot pressed Si_3N_4 was also impacted with SiC under the same particle size velocity conditions.

The results are discussed in terms of current erosion and impact models by considering particle size-velocity dependencies, appearance of the impact damage, and the basic properties and structure of the targets.

Under these erosion conditions, the four target materials exhibited widely different behavior not only in absolute amount of material removed, but also in mechanism of removal. The systems hot pressed Si_3N_4 -SiC particles and MgF_2 -quartz particles were characterized by plastic impact with associated lateral and radial crack formation, and erosion loss was proportional to particle mass and velocity to the fourth power. Erosion of hot pressed Si_3N_4 impacted with quartz particles was proportional to the first power of particle size and velocity, and loss occurred by minor chipping with no secondary crack formation. Erosion of glass bonded Al_2O_3 and reaction bonded Si_3N_4 did not show a consistent particle size-velocity dependence. The variation is related to the two phase structure of these materials. It was found that strength is not necessarily reduced for erosion conditions which produce appreciable material removal.

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1. INTRODUCTION

In recent years, considerable interest has been shown in the use of ceramics for high technology engineering components in such applications as gas turbine engines, bearings, heat exchangers, radomes and infrared transparent windows. All of these applications may involve impingement by solid particles. A knowledge of impact and erosion behavior is necessary before ceramics can be used with confidence in these systems. The materials discussed here, namely NC-132 hot pressed Si_3N_4 , NC-350 reaction bonded Si_3N_4 , Alsimag 614 glass bonded Al_2O_3 and Irtran 1 MgF_2 are either in current use or considered for potential use in one or more of the above applications.

To date, essentially two types of models have been proposed for impact or erosion of brittle materials (Refs. 1 and 2). The earlier models were based on elastic interaction and predicted that material removal occurs by the intersection of Hertzian ring cracks on the substrate surface. This process has been observed on several materials under static and low velocity impact conditions with relatively large spherical particles (for example, see Ref. 3). More recent analysis has treated static and dynamic plastic indentation which is characterized by plastic deformation of the contact area between the particle and the target, with radial cracks propagating outward from the contact zone, and with subsurface lateral cracks propagating outward on planes nearly parallel to the surface. The former are considered a source of strength degradation and the latter a potential source of material removal. These models are based on single impacts and were developed for isotropic materials under idealized conditions. One objective of this investigation was to assess the validity of these models to predict erosion of engineering ceramics by natural dust environments.

The experimental approach for this investigation has been to perform single impact and erosion (multiple particle impacts) tests in a controlled manner to simulate a service environment in the subsonic velocity regime. This approach advances the understanding of erosion mechanisms of engineering materials as well as providing data of direct value in application of these materials to engineering structures.

2. EXPERIMENTAL PROCEDURE

2.1 MATERIALS

The target materials exhibit a wide range of properties and structure. The properties considered pertinent to erosion response are listed in Table 1. Properties of the particles are also shown.

Angular, high purity quartz particles were used for most of the erosion testing. Quartz was chosen because in previous work on metallic erosion, it had been found to be the principal erosive component in natural dusts, i.e., the amount of erosion was directly proportional to the percentage of quartz in the natural dusts (Ref. 4). Six particle size ranges were used as

Table 1
Physical Properties of Target Materials and Particles

	Elastic Modulus GPa	Fracture Toughness MPa m ^{1/2}	Hardness ^a GPa	Acoustic Impedance Kgm ⁻² s ⁻¹ × 10 ⁷	Structure
Hot Pressed Si ₃ N ₄ (NC-132)	320	5	16	3.2	Pseudo single phase - 2μ grain size
Glass Bonded Al ₂ O ₃ (Alsimag 614)	324	3.2	12	4	2-phase - 4% glass, 96% Al ₂ O ₃ , 10μ grain size
Reaction Bonded Si ₃ N ₄ (NC-350)	170	2.2	3	2	Multiphase - Si ₃ N ₄ + porosity + Si + SiO ₂ (surface)
Hot Pressed MgF ₂ (Irtran I)	170	1	6	3.2	Single phase - ~2μ grain size
Natural Quartz	95	~0.7	~6	1.6	
SiC	420	~3	~23		

^aThe hardnesses are the quasi-static Vickers hardness in the macro indentation load independent regime.

follows: less than 30, 44-53, 62-74, 105-125, 250-297 and 350-420μm. These size ranges were chosen to be representative of airborne dust and to provide significant particle mass differences of at least 1/2 order of magnitude.

Angular silicon carbide particles of the following size ranges: less than 24, 25-85, 50-165, 203-495, 356-813, and 660-1346μm were supplied by Bendix Abrasives Division. The primary reason for selecting SiC as the second particulate was because of its much higher elastic modulus and hardness compared with quartz. However, the results of the SiC particle tests have direct practical value because it is used as an additive in aircraft carrier antiskid decking.

2.2 EROSION TESTING

Erosion tests were performed with a stationary target impacted by particles accelerated in an air stream. Particles are injected into the stream three meters from the target to provide sufficient distance for acceleration. High pressure, filtered and chemically dried air is used for the particle carrier gas. The carrier air velocity is measured using standard Pitot tube techniques. The air velocity variation across the 0.95 cm diameter nozzle is less than five percent and velocity is varied between 15 and 343 m/sec to achieve the desired particle velocity.

Particle velocity is measured using the rotating double disc technique described in Reference 5. Comparison with calculated velocities based on two phase flow theory is good (Ref. 6). Five velocities for each particle size range were used to establish erosion rates.

All erosion tests were performed at 90 degree impingement angle at ambient temperature. Perpendicular impingement is at or near that for maximum erosion of brittle materials. The number of particles used per test was varied from a few particles (to examine single particle impacts) to as many as 10^8 (to insure initiation of uniform erosion and to avoid incubation effects) over a 0.71cm^2 target area. The particles are fed into the gas stream using a precision feeder at a sufficiently low concentration such that particle interactions in the carrier gas stream or on the target surface are negligible. For the longtime-large number of particle tests, the specimens were weighed at specific time intervals to assess any changes in erosion with number of impacts. A detailed description of the erosion test apparatus is given in Reference 6.

2.3 STRENGTH MEASUREMENTS

Strength was determined in three point bending for both forms of Si_3N_4 and the Al_2O_3 . Baseline values were obtained on as-received material. To determine a strength loss threshold, strength of eroded specimens was measured over a wide range of test conditions from preweight loss to deeply eroded specimens. Each specimen was visually examined to insure that failure originated in the area subjected to particle bombardment. The eroded areas which are circular of 1 cm diameter were located in the center of the 2.5 or 1.25 cm specimens to insure no extraneous edge effects on strength measurements.

2.4 EXAMINATION OF ERODED SURFACES

The eroded surfaces were examined both optically, and by replica transmission and scanning electron microscopy. The progression of impact or erosion events was monitored by examining the specimens after various number of impacts for several particle size - velocity combinations ranging from single particle impacts to deeply eroded surfaces. The surfaces were also examined in cross section to assess the nature of subsurface damage.

3. EXPERIMENTAL RESULTS AND DISCUSSION

This section is separated into four parts. Initially, the erosion dependence on particle size and velocity will be presented for the 5 target-particle combinations, followed by discussion and examples of the type of impact damage that occurs. Effect of erosion on strength will then be presented, and in the final section a comparison between erosion behavior in terms of impact models and material properties and structure will be made.

3.1 EROSION FUNCTION OF PARTICLE SIZE AND VELOCITY

The customary method of reporting engineering erosion data is by plotting erosion weight loss versus some function of the erosive environment (i.e., particle size, velocity, weight of dust ingested, etc.). Under the same erosion conditions this approach provides a relative ranking of erosion response of the various target materials. Additionally, the phenomenological models proposed for erosion response include dependent functions for particle size and velocity, so that a knowledge of the particle size and velocity dependency of erosion behavior is an important first step to understanding the actual erosion mechanism.

Erosion weight loss as a function of particle mass and velocity was determined for conditions involving millions of particle impacts. This large number of impacts was necessary to insure that uniform erosion is occurring, and that effects from starting surface condition were minimized. In the case of reaction bonded Si_3N_4 (RB Si_3N_4) the as-fired surface layer was removed prior to erosion testing because it was found to erode at a much more rapid rate than the parent material. Weight loss per particle was plotted versus particle radius (log-log plot) and the slopes at constant velocity were measured to determine the particle size exponent. A similar plot of weight loss per particle versus particle velocity yielded the particle velocity exponent at constant particle size.

Two distinct relationships were observed. For the systems, RB Si_3N_4 and MgF_2 impacted with quartz particles and hot pressed Si_3N_4 (HP Si_3N_4) impacted with SiC particles, erosion is proportional to the fourth power of both particle size and velocity. The results are shown in Figure 1 for the three systems. The data is shown as volume loss to give a more meaningful comparison for the different density targets. The relationship between R^4V^4 and erosion is valid over 8 orders of magnitude for MgF_2 -Quartz and HP Si_3N_4 -SiC, which corresponds to 6 particle size ranges and 5 velocities between 24 and 285 mps for each particle size. The relationship also applies for RB Si_3N_4 -Quartz for all particle sizes at higher velocities. These results suggest that a single mechanism is controlling erosion under these test conditions. The data points associated with arrows correspond to weight increases that occurred after low velocity erosion tests on RB Si_3N_4 . Since RB Si_3N_4 is an inherently porous (~25% porosity) material, it is probable that the weight increases correspond to embedding of the quartz particles in the target. This will be discussed later in more detail.

For a given erosion condition with quartz particles, the amount of MgF_2 lost per impact is approximately 1-1/2 orders of magnitude greater than for RB Si_3N_4 . HP Si_3N_4 impacted with SiC particles is intermediate between the two.

The system HP Si_3N_4 impacted with quartz particles showed a significantly different erosion dependency on particle size and velocity than the systems previously discussed. Erosion is proportional to particle size to the third power and velocity to the first power which is a measure of particle momentum

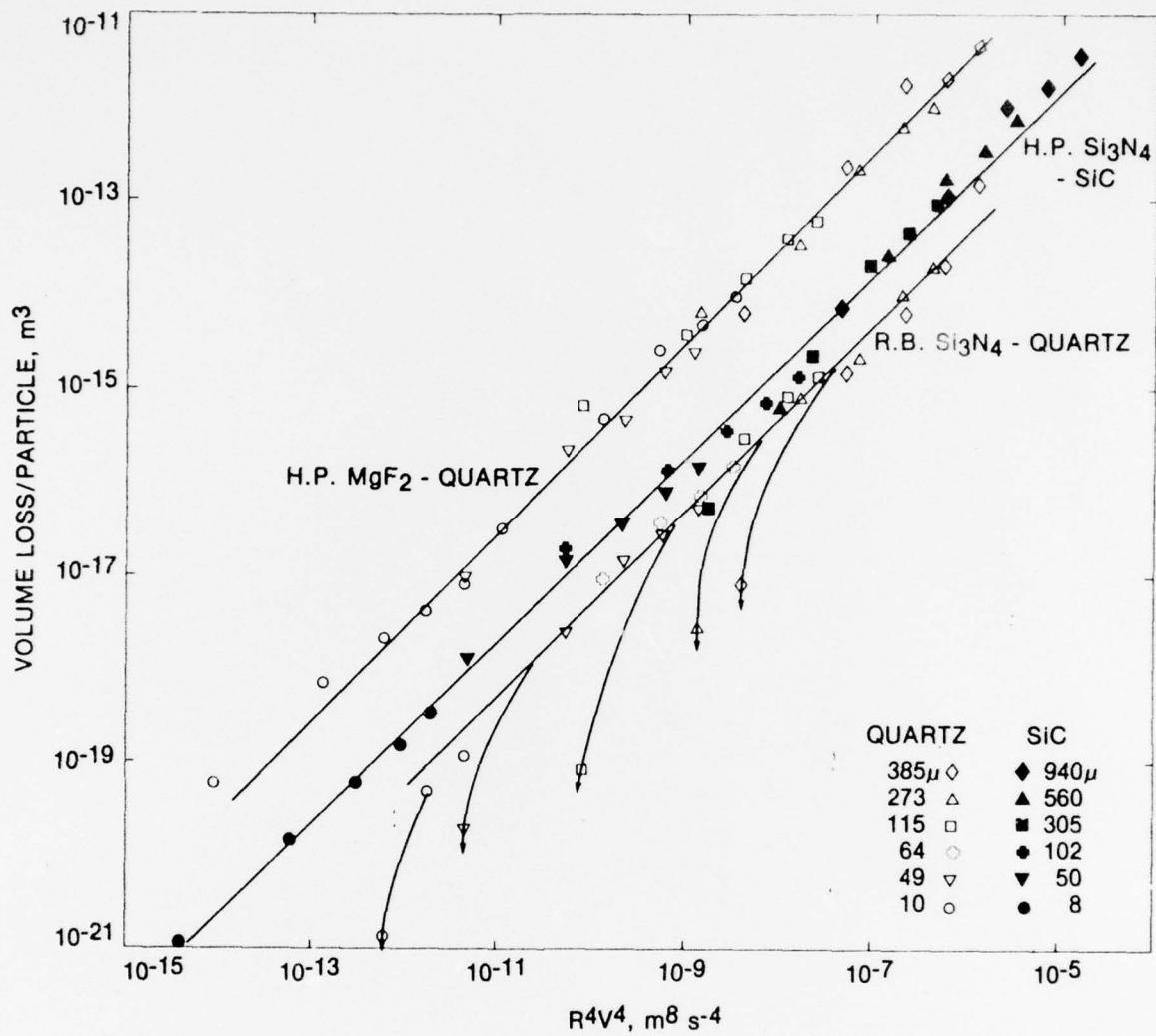


Figure 1. Erosion Versus Particle Size and Velocity to the Fourth Power (velocity varied between 40 and 285 m/sec)

(particle mass times velocity). The results for this system are shown in Figure 2 plotted as volume loss per particle versus a measure of particle momentum. The results are valid over the entire range of particle size - velocity combinations investigated. Deviations from the straight line relationship occur as the volume or weight loss threshold is approached for 115 and 273 μ particles. The results for this system indicate that a single mechanism or erosion process is operative within this range of test conditions and that the mechanism of erosion of HP Si₃N₄ by quartz particles is significantly different than the other three target - particle systems which showed a R⁴V⁴ erosion dependence. For equivalent particle size - velocity tests on HP Si₃N₄, the amount of material lost with SiC impacts was 2 orders of magnitude greater than for quartz impacts (highest velocity tests).

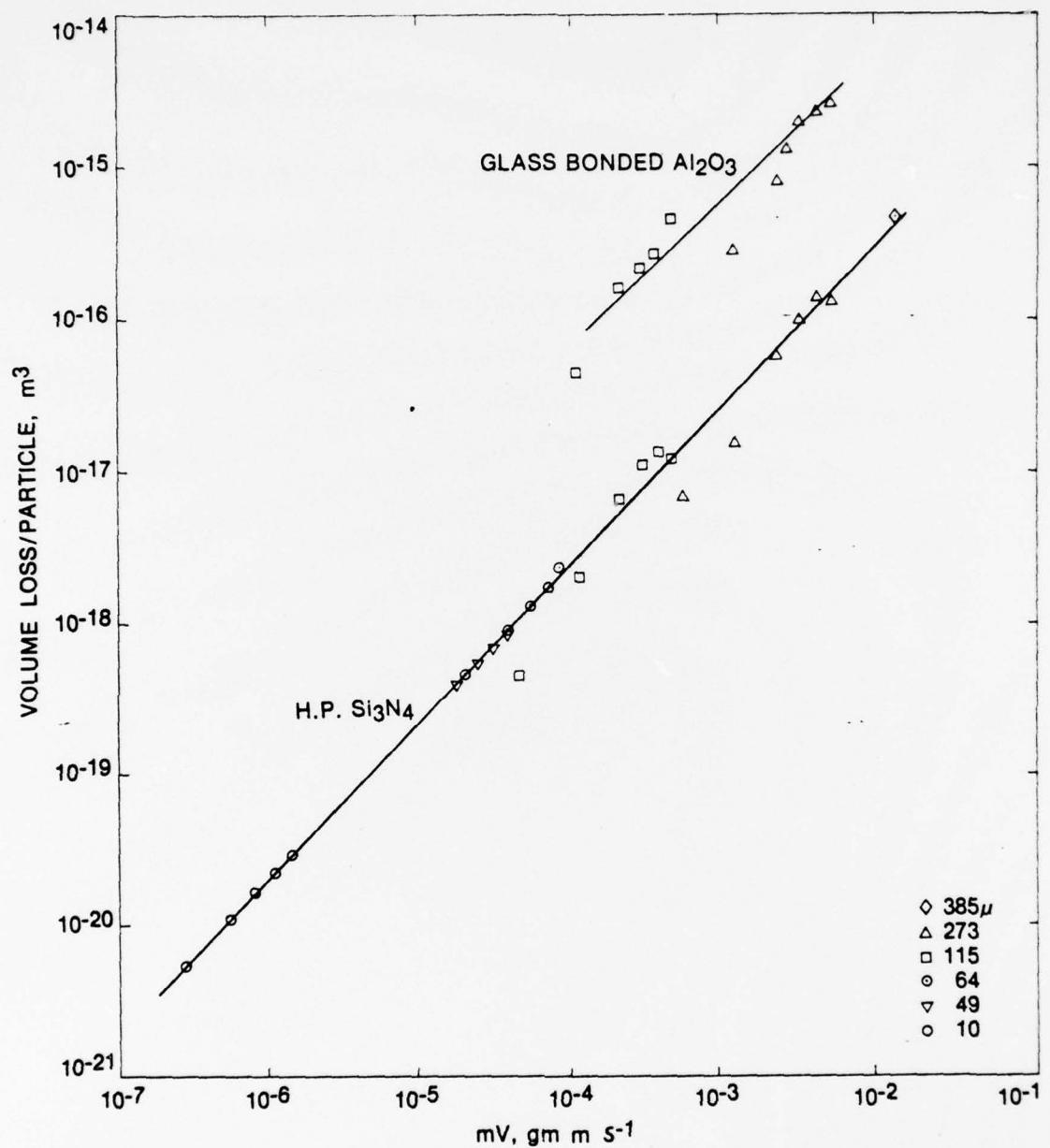


Figure 2. Erosion by Quartz Versus a Measure of Particle Momentum (velocity varied between 40 and 285 m/sec)

Glass bonded Al₂O₃ (GB Al₂O₃) impacted with quartz particles did not exhibit a consistent particle size - velocity dependence. Erosion was proportional to particle radius cubed over the entire range of particle sizes, but the velocity exponent varied between 3 for the smaller particles and one for the larger particles at higher velocities. The test data for erosion with the larger particles is also shown in Figure 2. Apparently, the rate controlling erosion mechanism or process is changing for these test conditions. This change is believed to be related to the 2-phase structure of the target and will be discussed further in the following section.

3.2 EXAMINATION OF ERODED SURFACES

Surfaces were examined for a range of erosion conditions varying between single particle impacts and erosion to a depth of several grain diameters for the various particle size velocity test conditions. Generally, the heavily eroded surfaces were sufficiently damaged that little information was provided concerning erosion mechanisms or processes. However, single particle impacts have provided insight into the process of erosion for these target materials. Figure 3 shows examples of single particle impacts for those target-particle combinations which exhibited a consistent particle-size velocity dependence. The types of impact are typical for each system although magnitude varied with particle size and velocity. The contact areas estimated for purely elastic impact by a spherical particle are given to illustrate variations in damage magnitude. HP Si₃N₄ impacted with SiC particles and MgF₂ impacted with quartz particles (Fig. 3a and 3b) both exhibited plastic impact accompanied by radial and lateral crack formation. The radial cracks extend outward from the particle contact zone and are generally perpendicular to the surface. Lateral cracks also extend from the contact zone but are subsurface and approximately parallel to the surface. This type of impact damage has recently been observed in other systems (Ref. 2). Material removal occurs by loss of a portion or all of the laterally cracked regions. The dark semicircular regions in Figures 3a and 3b correspond to laterally cracked material which has been removed during impact. Also shown in Figure 3a by polarized light reflecting conditions are the lateral cracks which have not resulted in material loss. Since HP Si₃N₄ is opaque, subsurface cracks are not observable by the polarized light technique. For these two systems, the extent of damage or material loss is much larger than the particle contact area.

Figure 3c shows a quartz particle impact on RB Si₃N₄. The impacts in this system are characterized by relatively deep pits with no apparent evidence of secondary cracks intersecting the surface. The approximate calculated contact radius ($\sim 35\mu$) is of the same order as the pit. The appearance of the pit is one of plastic impact. However, since this material is inherently porous ($\sim 25\%$), the phenomena could be one of crushing rather than plastic deformation. This porosity might also be expected to arrest crack propagation which would occur in a fully dense material. The type of impact

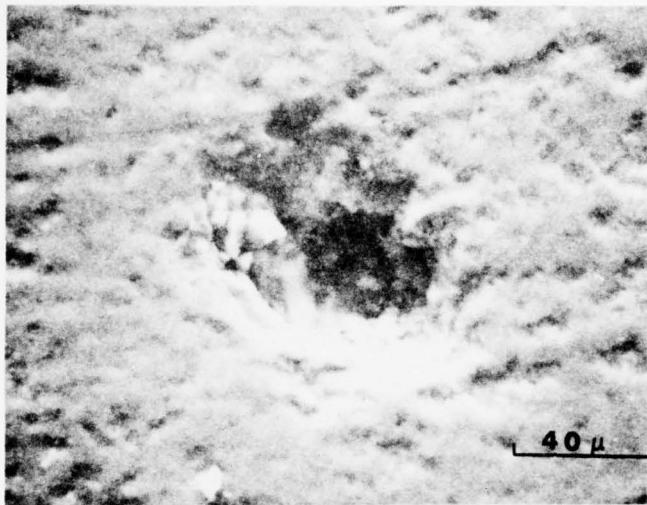


a. Hot Pressed MgF_2 Impacted
With 273μ Quartz
188 m/sec Velocity
Contact Radius $\sim 27\mu$
(3 impacts)



b. Hot Pressed Si_3N_4 Impacted
With 305μ SiC
176 m/sec Velocity
Contact Radius $\sim 22\mu$

Figure 3. Single Particle Impacts for Systems Which Exhibit Consistent
Particle Size-Velocity Functions With Erosion (Sheet 1 of 2)



c. Reaction Bonded Si_3N_4
Impacted With 385μ Quartz
174 m/sec
Contact Radius $\sim 35\mu$



d. Hot Pressed Si_3N_4 Impacted
With 273μ Quartz
188 m/sec Velocity
Contact Radius $\sim 25\mu$

Figure 3. Single Particle Impacts for Systems Which Exhibit Consistent
Particle Size-Velocity Functions With Erosion (Sheet 2 of 2)

damage shown in Figure 3c is characteristic of the RB Si_3N_4 quartz particle system except for the smallest particles (10μ) where single impacts could not be differentiated from the inherent porosity. Examination in cross section revealed radial cracking beneath the contact area that propagated through the pores.

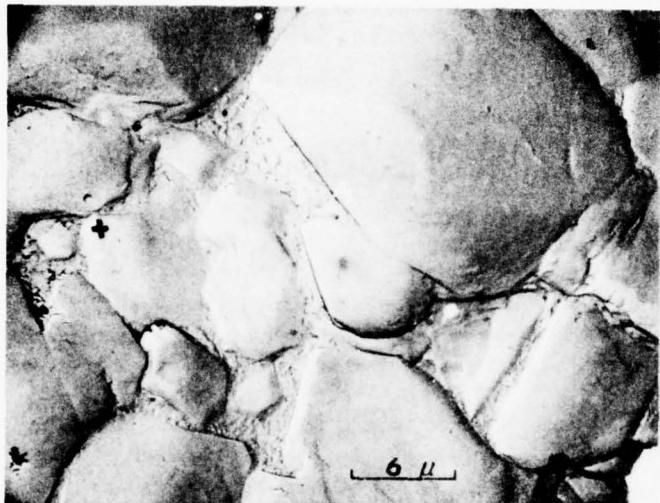
The type and magnitude of damage produced by quartz particles on HP Si_3N_4 is entirely different from that of the three systems previously discussed. An example is shown in Figure 3d. The chip which has been removed ($\sim 4\mu$ dia) is quite small compared with the approximate calculated contact diameter. For a corner oriented impact with an angular particle the contact area would be smaller and the stresses would be greater.

An estimate of the volume of material removed per impact can be made from the weight loss data on heavily eroded surfaces (data from Figs. 1 and 2). For all of the systems shown in Figure 3, the volume removed per impact for heavily eroded surfaces is within a factor of 2 of that measured from single particle impacts. Considering the statistical nature of the calculations and the test procedure, this shows quite good agreement between single particle impacts and bulk erosion and suggests that secondary cracking and residual erosion damage may not play a significant role in the erosion process under these test conditions.

The discussion to this point in this section has concerned the target-particle systems which exhibited a uniform particle size-velocity relationship over the entire range of test conditions (except low velocity impacts on RB Si_3N_4). The system GB Al_2O_3 -Quartz particles did not show a consistent relationship. The structure of this material is characterized by Al_2O_3 grains ($\sim 10\mu$ diameter) surrounded by a glassy intergranular phase. The glass can be as wide as 4μ in the vicinity of grain boundary triple points. It was found that single impact damage for 10μ impacts produced plastic flow in the grain boundary phase and minimal damage to the Al_2O_3 grains. The erosion process occurred by flow and removal of the glass to the extent that entire grains were lost. An example of a surface heavily eroded by 10μ quartz is shown in Figure 4. The depth of erosion corresponds to ~ 4 grain diameters. As can be seen, little damage has been sustained by the Al_2O_3 grains. As the particle size was increased, chipping of the Al_2O_3 grains occurred in addition to flow of the glass. This chipping is similar in appearance to that which occurs in the HP Si_3N_4 -Quartz system.

Hertzian type cone or ring cracking was not observed for any of these systems under these test conditions.

Examination of the impacted and eroded surfaces has shown that more than one type of impact occurs for this group of target materials under these test conditions. A discussion of the reasons for this variation based on properties and structure of the targets and particles will be given in the General Discussion section.



285 m/sec Velocity
Contact Radius $\sim 2\mu$
Depth of Erosion $\sim 50\mu$
6000 Particle Impacts per
Single Particle Contact Area

Figure 4. Glass Bonded Al_2O_3 Impacted With 10μ Quartz

3.3 STRENGTH OF ERODED SPECIMENS

Strength after heavy erosion was determined for HP Si_3N_4 , RB Si_3N_4 and GB Al_2O_3 all impacted with quartz particles (same test conditions used to develop Figs. 1 and 2 plus specimens eroded at particle size-velocity conditions below the weight loss thresholds). It was originally expected from Hertzian type considerations that a strength decrease would occur for impact conditions below those needed for actual weight loss, i.e., cracking would occur, but be insufficient to produce material loss. Above the weight loss threshold, strengths were expected to decrease.

The strength results are shown in Table 2. The baseline strengths plus one standard deviation are given for each target material. An estimate of the critical flaw size is also shown. This was calculated from fracture toughness values given in Table 1 using the relationship for three point bend specimens (Ref. 7). The strengths after erosion are shown as a function of the base line strengths by giving the percentages which have strengths greater than +1, within ± 1 , and less than -1 standard deviation of the noneroded material strength.

HP Si_3N_4 and the GB Al_2O_3 did not exhibit a strength decrease under these test conditions for erosion depths up to 37μ . For HP Si_3N_4 this depth corresponds to 3×10^8 particle impacts or 600 grams of dust on a 0.71 cm^2 area. There was a trend toward strength increase (fracture stress for 50% of the specimens was greater than one standard deviation above the baseline strength) which indicates a "polishing" phenomenon may be occurring. No specimens failed at a stress lower than the one standard deviation band. The erosion depth is approximately three times the estimated critical flaw size

Table 2
Strength Distribution of Eroded Specimens Relative to Standard Strength

	Pre-Erosion Baseline		Post-Erosion Strength Distribution, %					
	$\bar{\sigma}_F \pm 1$ Std. Dev. MPa	% Critical Flaw Size, μ	No. Tests	Max. Erosion Depth, μ	Above +1 Std. Dev.	Within ± 1 Std. Dev.	Below -1 Std. Dev. .	
HP Si_3N_4	798 ± 128	11	25	31	50	50	0	
GB Al_2O_3	252 ± 21	43	51	35	41	43	16	
RB Si_3N_4	278 ± 10	17	17	360	6	12	82 ^a	

^aMinimum Strength = 81 MPa Corresponding to Critical Flaw Size of $\sim 200\mu$

which indicates that under these test conditions, the effective flaw size produced by erosion is no larger than pre-existing flaws characteristic of the "standard" machined surface. (Because large numbers of flaws are introduced by erosion or machining, the effective stress concentration will depend on both size and spacing of flaws. In contrast, critical flaw sizes were calculated for isolated flaws.) In confirmation, examination of the eroded surfaces in cross section did not reveal apparent subsurface damage, i.e., the structure under the eroded area was indistinguishable from that below the as-machined surface and subsurface cracking was not apparent.

The trend toward a strength increase is not as conclusive with GB Al_2O_3 as with HP Si_3N_4 since 16% of the specimens failed at stresses below the one standard deviation range. Furthermore, although the maximum depth of erosion ($\sim 35\mu$) is approximately four grain diameters, this also corresponds to the estimated critical flaw size of the baseline material. However, the results indicate that erosion under these test conditions do not produce flaws greater than those inherent to the as-sintered surface.

RB Si_3N_4 did exhibit a marked strength decrease under these erosion test conditions. Strength decreased rapidly for the first 100μ erosion depth and remained essentially constant at 100μ up to the maximum depth tested (350μ). The tests were performed in the as-fired surface condition where the surface oxide layer is $\sim 100\mu$ deep. The lowest strength (81 MPa) corresponds to an increase in estimated critical flaw size over the baseline material of one order of magnitude (17μ compared with 200μ). The depth of subsurface cracks perpendicular to the surface is also $\sim 200\mu$ for this erosion condition.

According to the model for plastic impact response, the strength of the target (σ_F) should be dominated by radial crack formation by the following relation:

$$\sigma_F \propto \frac{K^{1.4}}{\sqrt{C} R^{0.8}}$$

Ref. 2

where K_C is target fracture toughness, V is particle velocity and R is particle radius. A log-log plot of strength versus $(V^{0.5}R^{0.8})^{-1}$ for reaction bonded Si_3N_4 is shown in Figure 5. The data falls roughly into two groups which both have slopes close to one on a log-log basis. The major difference between these two groups is particle size and depth of erosion. The depth of erosion for these particular specimens subjected to 10μ particle bombardment was still within the surface layer which has different material properties and composition than the bulk material. Furthermore, the type of damage and hence flaw type apparently differs from that produced with the larger particles. Subsurface cracking is minor compared with that produced with larger particles, and embedding of the 10μ particles was observed.

Since quartz is the most erosive constituent of natural dusts and the particle size-velocity conditions are typical of airborne dust, these results have direct practical significance, and indicate that a strength decrease does not necessarily occur for erosion conditions which produce appreciable material removal (HP Si_3N_4 and Alsimag 614 glass bonded Al_2O_3). This would be expected to apply as long as the load bearing volume is not reduced significantly.

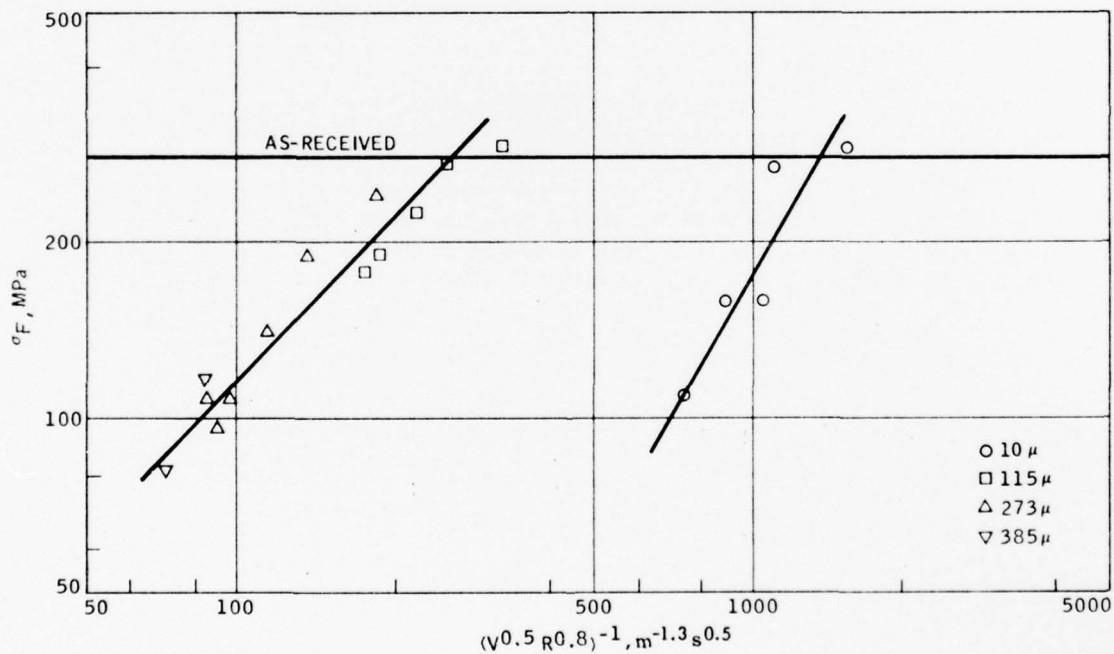


Figure 5. Strength Versus Function of Predicted Radial Crack Formation for Reaction Sintered Si_3N_4 Eroded by Quartz

3.4 GENERAL DISCUSSION

In this section, a comparison will be made between the erosion behavior of the various target materials, and the results will be discussed in terms of impact models and material properties and structure.

Four "engineering" ceramics have been subjected to erosion conditions considered characteristic of a natural dust environment. Under these conditions, the erosion response differs markedly (Figures 1 and 2), and for a given condition, there is approximately one order of magnitude difference in volume loss between each of the targets. There is also considerable variation in material properties and structure between the targets. Table 1 lists the physical properties considered relevant to erosion response. Of the properties listed, fracture toughness is the only property which varied consistently with erosion. The other properties (elastic modulus, hardness and acoustic impedance) would predict either less separation in erosion loss or a different ranking of erosion resistance. Furthermore the particle properties can also have a strong effect on erosion as evidenced by the fact that erosion of HP Si_3N_4 by SiC particles is approximately 2 orders of magnitude greater than with quartz particles for equivalent particle size-velocity test conditions, and the type of impact damage is entirely different.

To date, essentially two types of models have been proposed for solid particle impact or erosion at subsonic velocities. The earlier model was based on Hertzian type purely elastic interaction where material removal occurs by the intersection of ring cracks on the substrate surface. This model was expanded to include Weibull statistics whereby erosion was a function of the distribution and size of flaws under the particle contact area (Ref. 1). This relationship predicted that

$$E \propto R^4 V^{2.7}$$

(where E is volume removed per particle) for a Weibull modulus greater than 12 and impact with angular particles (Ref. 1). This velocity dependence was not observed for these test conditions, nor was ring cracking.

The second model has treated static, and more recently dynamic, plastic indentation which is characterized by plastic deformation of the contact area between particle and target, radial cracks propagating outward from the contact zone, and lateral cracks that initiate beneath the contact zone and propagate between the radial cracks on planes nearly parallel to the surface. It has been observed for a variety of single phase targets impacted with relatively incompressible projectiles (impact conditions above plastic impact threshold) that the length of radial cracks, and the depth of lateral cracks, for single impacts, followed the relationships predicted by the model (Ref. 2).

If it is assumed that maximum erosion loss per impact is proportional to the volume encompassed by the lateral cracks, then the plastic impact model predicts

$$E \approx V^{2.5} R^4 \left(\frac{\rho_p}{H K_c^6} \right)^{0.25} f(M)$$

where ρ_p is particle density, H is target hardness, and $f(M)$ is the fraction of the volume encompassed by the lateral cracks that is actually removed and is considered a material dependent variable. Since the experimental erosion data for this type of impact (HP Si_3N_4 impacted with SiC, and MgF_2 impacted with quartz) revealed both a radius and velocity dependence to the fourth power, the term $f(M)$ would have to include a velocity dependence, to obtain conformance with experimental data. This would require that the formation of lateral cracks as well as the volume of laterally cracked material actually removed be velocity dependent.

A log-log plot of $E/V^{2.5} R^4$ versus $\rho_p/H K_c^6$ for the four target materials shows a slope of 0.25 between the 2 target-particle conditions which exhibited typical plastic impact response (Fig. 6). The erosion data in Figure 6 is an average of all of the particle-size velocity test conditions \pm one standard deviation. However, a plot of $E/R^4 V^4$ versus $\rho_p/H K_c^6$ using all of the erosion data (much smaller data band than for $E/R^4 V^{2.5}$) also exhibited a slope of 0.25 between HP Si_3N_4 -SiC and MgF_2 -Quartz.

Erosion for the other target-particle combinations falls below that expected from the plastic impact model although the relative rank is consistent with the model. In the case of HP Si_3N_4 impacted with quartz particles, this is expected since the impact conditions are apparently below the plastic impact fracture threshold (material removal occurs by minor chipping with no apparent secondary crack formation). However, GB Al_2O_3 and RB Si_3N_4 did exhibit plastic deformation.

The model assumes an isotropic material. For the purposes of this investigation, both MgF_2 and HP Si_3N_4 can be assumed to be single phase materials (HP Si_3N_4 contains a minor grain boundary phase, but the 2μ grain size would average this effect over the particle contact area). However, GB Al_2O_3 and RB Si_3N_4 are not single phase materials. RB Si_3N_4 contains $\sim 25\%$ porosity, and although the impact craters had the appearance of plastic impact, it is possible that crushing is occurring under the contact area. Support for this hypothesis is given by the fact that particles embedded in the surface at low velocities for all particle sizes. Also, the lack of secondary cracking outside of the contact area may be related to the crack blunting ability of the pores. Erosion of GB Al_2O_3 is a two step process which is directly attributable to the two phase nature of the structure.

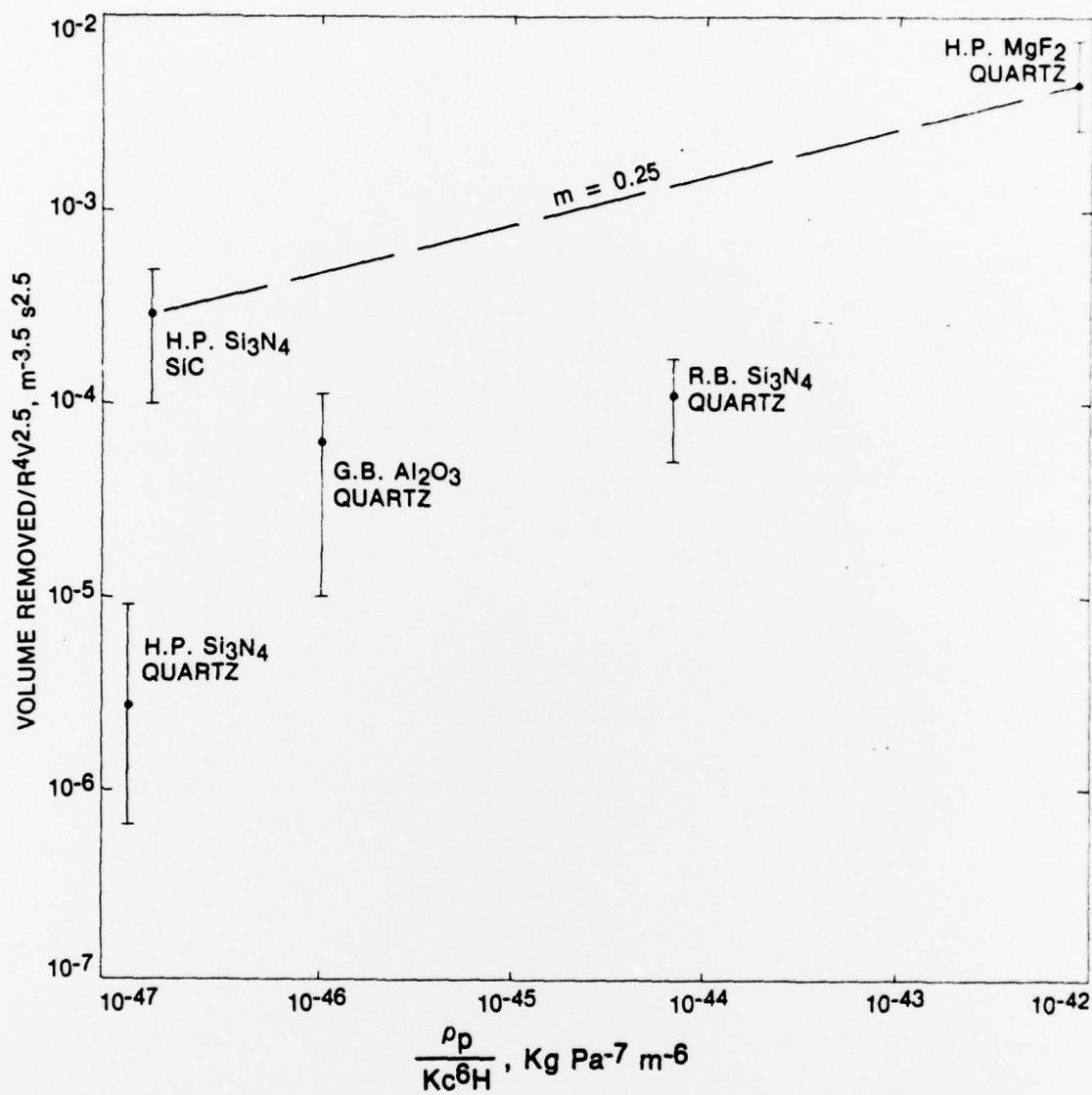


Figure 6. Relationship Between Erosion Data and Plastic Impact Model

The particle composition and properties obviously affect type of erosion response. However, from this investigation only general comments can be made concerning the influence of particle type on erosion of a given target. It appears that for plastic impact response, the particle mechanical properties must closely approach or exceed those of the target. For impact and erosion conditions where the particle properties are appreciably lower than the target, the erosion process is less efficient.

4. CONCLUSIONS

Four ceramic materials with widely different properties and structure were eroded under conditions which simulate a service dust environment i.e., 10-385 μ natural quartz particles at subsonic velocities. Additionally, one of the targets (HP Si₃N₄) was impacted with SiC particles over the same particle-size velocity conditions. The following particle size-velocity relationships for erosion were observed:

$$\begin{aligned} E \propto R^4 V^4 & \quad \text{MgF}_2\text{-Quartz, RB Si}_3\text{N}_4\text{-Quartz} \\ & \quad \text{HP Si}_3\text{N}_4\text{-SiC} \\ E \propto R^3 V & \quad \text{HP Si}_3\text{N}_4\text{-Quartz} \\ E \propto R^3 V^{1-3} & \quad \text{GB Al}_2\text{O}_3 \end{aligned}$$

Single particle damage for the systems HP Si₃N₄ impacted with SiC and MgF₂ impacted with quartz was characterized by plastic impact and radial and lateral cracks propagating from the contact area. The diameter of material removed could be as much as 3 times the contact diameter (corresponds to a portion of lateral crack formation). The diameter of material removed for RB Si₃N₄ was essentially the same as the estimated contact diameter and either plastic deformation or crushing occurred. Erosion of HP Si₃N₄ impacted with quartz occurs by minor chipping which is an order of magnitude less than the estimated contact radius. In the latter two cases, secondary cracking was not observed on the surface. However, RB Si₃N₄ exhibited extensive subsurface radial cracks under the contact area. Erosion on glass bonded Al₂O₃ is a two-step process involving plastic deformation of the glass plus chipping of Al₂O₃ grains.

Based on these results, erosion is a function of particle radius and velocity to the fourth power for a plastic impact response. For a single phase material (HP Si₃N₄, MgF₂) erosion appears to follow the function of fracture toughness and hardness predicted by the model for plastic impact. However, for multiphase materials, structure also influenced erosion response. Below the plastic impact threshold, erosion is a much lesser function of velocity, and elastic ring cracking was not observed. The plastic impact threshold was more a function of target particle type than of particle size and velocity.

Strength measurements after erosion with quartz revealed that strength is not necessarily reduced after conditions which produce significant material removal. (HP Si₃N₄ and GB Al₂O₃ did not show a strength decrease, while RB Si₃N₄ exhibited a major strength decrease.)

It is concluded that more than one mechanism of erosion exists under "natural" dust environments, and that the mechanisms are dependent not only on target physical properties, but also to a large extent on structure.

REFERENCES

1. Oh, H.L., Oh, K.D.L., Vaidyanathan, S. and Finnie, I., *On the Shaping of Brittle Solids by Erosion and Ultrasonic Cutting*, NBS Special Publication 348, 1972, p. 119.
2. Evans, A.G., Gulden, M.E., Eggum, G.E. and Rosenblatt, M., "Impact Damage in Brittle Materials in the Plastic Response Regime," Contract N00014-75-C-0069, Report No. SC5023.9TR, Rockwell International Science Center, Thousand Oaks, CA, 1976.
3. Adler, W.F., "Analysis of Multiple Particle Impacts on Brittle Materials, AFML Tech. Report 74-210, Wright-Patterson Air Force Base, OH, 1974.
4. Smeltzer, C.E., Gulden, M.E., McElmury, S.S. and Compton, W.A., "Mechanisms of Sand and Dust Erosion in Gas Turbine Engines," USAAVLABS Tech. Report 70-36, Fort Eustis, VA, 1970.
5. Ruff, A.W. and Ives, L.K., *Wear*, Vol 35, 1975, p. 195.
6. Gulden, M.E. and Metcalfe, A.G., "Study of Erosion Mechanisms of Engineering Ceramics," Solar Report RDR 1778-4, Solar Turbines International, San Diego, CA, Contract N00014-73-C-0401, 1976.
7. Evans, A.G. in *Fracture Mechanics of Ceramics 1*, R.C. Bradt, D.P.H. Hasselman and F.F. Lange, Ed., Plenum Press, NY, 1974, p. 17.

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Four "engineering" ceramics were subjected to impact (single particle) and erosion (multiple impacts) under conditions which simulate a natural dust environment in the subsonic velocity regime. The target materials are hot pressed Si ₃ N ₄ , reaction bonded Si ₃ N ₄ , glass bonded Al ₂ O ₃ and hot pressed MgF ₂ . Tests were performed with 6 narrow size ranges of natural quartz		

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between 10 and 385¹ average, and 5 velocities for each particle size. Hot pressed Si₃N₄ was also impacted with SiC under the same particle size velocity conditions.

The results are discussed in terms of current erosion and impact models by considering particle size-velocity dependencies, appearance of the impact damage, and the basic properties and structure of the targets.

Under these erosion conditions, the four target materials exhibited widely different behavior not only in absolute amount of material removed, but also in mechanism of removal. The systems hot pressed Si₃N₄-SiC particles and MgF₂-quartz particles were characterized by plastic impact with associated lateral and radial crack formation, and erosion loss was proportional to particle mass and velocity to the fourth power. Erosion of hot pressed Si₃N₄ impacted with quartz particles was proportional to the first power of particle size and velocity, and loss occurred by minor chipping with no secondary crack formation. Erosion of glass bonded Al₂O₃ and reaction bonded Si₃N₄ did not show a consistent particle size-velocity dependence. The variation is related to the two phase structure of these materials. It was found that strength is not necessarily reduced for erosion conditions which produce appreciable material removal.

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